NEW OPPORTUNITY

REGULATORY AFFAIRS MANAGER

ZUG AREA, SWITZERLAND

Do you enjoy overseeing the full life-cycle of medical devices? Do you feel limited in your tasks and your scope? Are you an experienced Regulatory Affairs professional in the medical device industry?

Yes? We have a fabulous opportunity for YOU!

THE COMPANY

Renowned for its excellent work-life balance, cosmopolitan cities and stunning natural beauty, Switzerland also hosts one of the biggest MedTech hubs in Europe. And, with its central location, travelling to anywhere in Europe only takes a couple of hours!

This is a very unique chance to join a family owned company of 2000+ people, at the company's HQ in Zug. You'll be at the centre of where all the decisions are made, meet senior leadership in the corridors, and have lots of face to face contact with your partners.

THE OPPORTUNITY

In this role as Regulatory Affairs Manager, you'll join a close knit team that is responsible for the regulatory management of medical devices, globally. With a presence in over 80 countries across EMEA, North America, LATAM and Asia, you'll have exposure to global markets and get to learn about the regulatory environment in each of those countries.

This is a really varied role where you'll have the chance to work on new product development projects AND oversee the whole lifecycle of a device, from concept/development to market.

You'll also support the MDR transition!

REQUIREMENTS

- 5+ years of regulatory affairs experience in the **medical device** industry
- Fluent written and speaking skills in English

Interested in further conversation?

Please send your CV to kristina@elemed.eu to arrange a confidential career discussion.